Hereditary Cancer Genetic Test

What is the Hereditary Cancer Genetic Test?

The Hereditary Cancer Genetic Test analyzes 29 genes linked to inherited risks for common cancers, including breast, colorectal, uterine, ovarian, prostate, and more. This test can provide valuable insights to help you and your healthcare providers plan for early detection, preventive measures, and personalized care. Additionally, it provides insights into genetic ancestry and traits, allowing you to discover how your genes may influence your preferences, appearance, and other fun genetic facts.

What Does This Test Look For?



Hereditary Cancer Genetic Test: Analyzes 29 genes for genetic variants (also called mutations) that may increase your risk for common hereditary cancer types. These genes include:

APC, ATM, BAP1, BARD1, BMPR1A, BRCA1, BRCA2, BRIP1, CDH1, CDK4, CDKN2A (p14ARF), CDKN2A (p16INK4a), CHEK2, EPCAM, GREM1, MITF, MLH1, MSH2, MSH6, MUTYH, PALB2, PMS2, POLD1, POLE, PTEN, RAD51C, RAD51D, SMAD4, STK11, TP53

To learn more, see the Gene Tables on page 3.



Discovery: Explores insights about your genetic ancestry and learn about how your genes influence unique characteristics such as lactose intolerance, alcohol flush response, cilantro preference, and more.



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What are the possible results?

Something important is found

A meaningful genetic change is identified that could impact your health.

Nothing significant is found

No increased risk for the hereditary conditions were found in the genes tested.

How can these results impact my healthcare?

Earlier interventions, at every step: Insights into your hereditary cancer risk can guide your healthcare providers in recommending regular screenings or lifestyle changes that suit your needs.

Family health insights: Your results may also provide useful health information for family members, as some genetic risks can be shared among relatives.

What happens after I get my results?

Color's care team will ensure you understand the result and the next steps associated with the result to take action on your health. We recommend that you share your Color test results with your healthcare provider. This can help you and your provider create a personalized healthcare plan. If you don't have your own healthcare provider, Color's care team can help get you connected to one in your area.



Hereditary Cancer Genetic Test

| Gene | Hereditary Condition | Associated Cancers |
|--------|--|---|
| APC | Familial adenomatous polyposis (FAP), Attenuated FAP | Colon, thyroid, brain, stomach, small bowel |
| ATM | Hereditary breast and pancreatic cancer | Breast, pancreatic |
| BAP1 | BAP1 tumor predisposition syndrome | Melanoma, kidney, lung |
| BARD1 | Hereditary breast cancer | Breast |
| BMPR1A | Juvenile polyposis syndrome | Colon, stomach |
| BRCA1 | Hereditary breast and ovarian cancer syndrome | Breast, ovarian, pancreatic, prostate |
| BRCA2 | Hereditary breast and ovarian cancer syndrome | Breast, ovarian, melanoma, pancreatic, prostate |
| BRIP1 | Hereditary ovarian cancer | Ovarian |
| CDH1 | Hereditary diffuse gastric cancer | Breast, stomach |
| CDK4 | Familial atypical mole-malignant melanoma syndrome | Melanoma, pancreatic |
| CDKN2A | FAMMM, Melanoma and neural system tumor syndrome | Melanoma, pancreatic, nervous system |
| CHEK2 | Hereditary breast cancer | Breast, prostate |
| EPCAM | Lynch syndrome | Colon, uterine, ovarian, stomach, pancreatic, prostat |
| GREM1 | Hereditary mixed polyposis syndrome | Colon |
| MITF | Hereditary melanoma and kidney cancer | Melanoma, kidney |
| MLH1 | Lynch syndrome | Colon, uterine, ovarian, stomach, pancreatic, prostat |
| MSH2 | Lynch syndrome | Colon, uterine, ovarian, stomach, pancreatic, prostat |
| MSH6 | Lynch syndrome | Colon, uterine, ovarian, stomach, prostate |
| MUTYH | MUTYH-associated polyposis | Colon, uterine, ovarian, stomach, prostate |
| PALB2 | Hereditary breast and ovarian cancer | Breast, ovarian, pancreatic |
| POLD1 | Polymerase proofreading-associated polyposis | Colon |
| POLE | Polymerase proofreading-associated polyposis | Colon |
| PMS2 | Lynch syndrome | Colon, uterine, ovarian, stomach |
| PTEN | PTEN hamartoma tumor syndrome | Breast, thyroid, uterine, kidney, colon |
| RAD51C | Hereditary breast and ovarian cancer | Breast, ovarian |
| RAD51D | Hereditary breast and ovarian cancer | Breast, ovarian |
| SMAD4 | Juvenile polyposis syndrome | Colon, stomach |
| STK11 | Peutz-Jeghers syndrome | Breast, colon, stomach |
| TP53 | Li-Fraumeni syndrome | Breast, colon, brain, pancreatic, sarcoma |
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